



Method and Apparatus for Treating Obstructive Sleep Apnea Syndrome

References Disclosed

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U.S. Patent number **6,209,542 (Thornton)**;

U.S. Patent number 5,752,510 (Goldstein);

U.S. Patent number 6,012,455 (Goldstein);

"TAP/SAAMS" product literature (Airway Management, Inc.; Dallas, TX)

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U.S. Patent 4,782,832 (Trimble et al)

Background of the Invention

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Obstructive Sleep Apnea Syndrome (OSAS) is a serious medical condition that is difficult to treat easily and effectively. The condition is marked by a partial or complete closure of the upper airway during sleep.

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There are four basic therapeutic modalities in the treatment of OSAS: pneumatic splinting of the airway with positive airway pressure (PAP), airway orthotic (AO) which mechanically dilates the upper airway, surgery, and combination therapy which uses an airway orthotic plus positive airway pressure.

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PAP acts as a pneumatic splint which literally blows open the upper airway during sleep such that the tendency for the upper airway to collapse is either eliminated or reduced.

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There are a variety of surgeries to treat OSAS, some approaches work better than others. Since this is not the subject of this patent, nothing further will be said on this issue.

Combination therapy typically utilizes mechanical or physical tissue manipulation of the muscles of the upper airway via use of an AO plus pneumatic splinting of the airway. Alternatively, combination therapy may use an AO without mechanical dilation of the upper airway. When an AO is used without

5 mechanical dilation of the upper airway, the AO may be utilized to stabilize an application device for delivery of PAP with a non-claustrophobic interface and the active treatment then depends solely on pneumatic splinting of the upper airway.

Standard PAP therapy usually includes a nasal application device for delivery of

10 the positive airway pressure to the patient's nasal passages. There are different styles of nasal application devices. These include:

- a) Triangular-shaped nasal masks which form a seal surrounding the patient's nose on his facial surface,
- 15 b) Full-face masks that form a seal on the patient's face around both the nose and mouth,
- c) Nasal inserts commonly referred to as nasal pillows.

The nasal application device is connected to a Continuous Positive Airway

20 Pressure machine (CPAP) via tubing. These air pressure machines (CPAP) can be set to certain and particular pressure settings. Each patient will have a certain and particular pressure level of PAP that will dilate and pneumatically splint his upper airway. When utilized properly, this therapy is very effective in treating the patient with a nocturnally obstructed airway.

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Nasal CPAP is considered to be the gold standard in treatment of moderate to severe OSAS. However, this life saving therapy is fraught with significant compliance problems. Clinical studies indicate that compliance on nasal CPAP is less than 50%. Reasons for this non-compliance are numerous and include

30 problems with mask fit and resultant mask discomfort, mask leakage of PAP, claustrophobia, head gear discomfort, head strap discomfort, sleep position

(supine) confinement, mouth venting or leakage, chin strap confinement and discomfort, dermatitis, swallowing of air, pressure-related tolerance issues etc.

5 Airway Orthotics (AO) are oral appliances that are worn in the mouth at night for the purpose of treating OSAS. Historically, there are two basic types of airway orthotics: the tongue retention device (TRD) and the mandibular advancement device (MAD). For the purposes of this disclosure nothing more will be said about the TRD other than to note its existence.

10 Most mandibular advancement (MADs) devices mechanically dilate the upper airway by moving the lower jaw, or mandible, forward in controlled increments. The airway is dilated by a combination of: pulling the tongue base anteriorly away from the airway and holding it forward during sleep, tightening of the apneic patient's upper airway via anterior and superior movement of the hyoid bone
15 which stretches the infrahyoid muscles attached to the upper airway, innervation of key airway muscle groups such as the genioglossus, palatoglossus, and inferior bellies of the lateral pterygoid(s), and lateral stretching of the hypopharynx.

20 Mandibular advancement devices can either be fixed or adjustable. In general, adjustable MADs are preferred by most clinicians who treat OSAS because they may be titrated, or gradually adjusted, according to the patient's airway needs. These MAD devices incorporate a variety of different designs and schemes to move the lower jaw forward.

25 However, advancing of the mandible can result in unpredictable occlusal (bite) changes. Therefore, the clinician must weigh all possible outcome factors, both positive and negative, in formulating his treatment plan.

30 Mandibular advancing devices, while often helpful, can complicate issues for the patient undergoing "combination therapy." Complicating issues can compromise

the effectiveness of the therapy by resulting in decreased compliance or utilization. Therefore, there is a need to create simple and effective therapies for OSAS that minimize complications. Often, the effectiveness of nasal CPAP, airway orthotics, or combination therapy relates to patient compliance with the therapy. Simpler, less burdensome therapies will generally lead to higher compliance and resultant efficacy in treatment.

The objectives of this invention are to improve on the concept of combination therapy for OSAS based on several factors.

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- 1) Simplifying the use by the patient to increase compliance through minimization of necessary parts in design of the airway orthotic;
- 2) Accommodation of varying nare-to-nare and nose widths for use of nasal pillows and tubing;
- 15 3) Obturation of the oral cavity preventing mouth venting of PAP without the need for a chin strip or need to add a separate piece known as a "vent or saliva shield;"
- 4) Use of a 3mm minimum thickness of acrylic for PAP Tubing Retention Platform construction to improve strength, integrity and malleability to customize tubing angulation via application of heat. Usually this heat is applied by a micro torch;
- 20 5) Improving angulation of nasal tubing by closest preferred approximation antero-posteriorly to positions directly below the nares.
- 6) Minimization of occlusal changes in utilization of airway orthotics by building these oral appliances in a neutral-centric mandibular position with no mandibular advancement such that the mandible is in centric relation, centric occlusion or the like.
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Design of Airway Orthotic

This invention will introduce a new type of airway orthotic that will simplify insertion, usage, and increase effectiveness for the patient. This airway orthotic serves three primary functions:

- 5 a) to obturate or seal off the oral cavity preventing mouth breathing and/or mouth venting of positive airway pressure (PAP);
- b) to support application of PAP via nasal pillows directly from the obturator without use of any headgear, straps or chin support to stabilize the nasal seal during movements at night.
- 10 c) to hold the upper and lower jaws in a preferred position via use of an elastomeric material without the use of a hook, chin stabilizer, or strap.

15 **Method of Manufacture**

- 1) Individual impressions are made of the upper and lower teeth and master casts are poured in stone or plaster.
- 20 2) A bite registration is also recorded which documents the relationship of the mandible to the maxilla in three dimensions and is used by the laboratory to mount the upper and lower casts on an articulator. The bite registration position may be captured either:
 - 25 a) In neutral centric position, i.e., without mandibular protrusion/advancement; or,
 - b) In a protruded position referred to as mandibular advancement; and
 - c) May vary the caudal or vertical component to make room for the tongue and/or
 - 30 maximize patency of the airway.

3) The airway obturating orthotic is professionally manufactured in the laboratory and is composed of a hard acrylic exterior which is lined with an elastomeric material such that there is a snap fit when the upper and lower teeth are engaged into it. This snap fit via said elastomeric material holds the lower jaw in preferred
5 position without any additional means. Elastomeric materials flex like rubber when pressure is applied as in the case wherein teeth snapfit over the internal surfaces of the appliance.

The following passages will now compare the present invention with prior art
10 specifically referenced to illuminate useful and novel differences between the present invention and the prior art.

By way of reference I direct the reader to U.S. Patent 6,209,542, entitled
"Combination face mask and dental device for improved breathing during sleep."
15 Dr. Thornton describes an apparatus consisting of a face mask connected to an oral appliance. The adjustment means allows for positioning of the mask from a first to a second position. The patent describes an oral appliance that has separate upper and lower arch components.

20 Goldstein 5,752,510 and 6,012,055 teaches of a dual-arch mandibular advancement device that incorporates application of a nasal mask and nasal inserts.

By comparing the referenced prior art the reader will note fundamental
25 differences between the present invention and what is taught by Thornton and also Goldstein.

The airway orthotic or oral appliance in the present invention is *intentionally non-adjustable*. By utilizing a fixed/*non*-adjustable appliance the apnea patient avoids
30 the complication of:

- a) first inserting the upper arch and then the lower arch and:
- b) then hooking them together.

For certain patients, like those patients with a significant mental or developmental
5 impairment or impairment through aging, this joining together of individual
components can be challenging. This challenging problem alone can lead to
non-compliance and subsequent failure of the therapy.

10 With the present invention, the patient simply opens his mouth and bites down
with the upper and lower teeth simultaneously onto a one-piece, dual arch
orthotic in one clean motion. This fixed orthotic contains both the upper and
lower arches in one unit lined with an *elastomeric* material. This simplicity is
preferable because it will increase compliance on the therapy, which increases
overall effectiveness medically.

15 There is the problem referred to as *mouth venting of PAP*. This is where air is
blown into the nose and the patient inadvertently opens his mouth while asleep
and thereby permits air to be blown out of his mouth, via path of least resistance.
This occurs due to insufficient obturation or sealing-off of the oral cavity. This
20 results in inadequate requisite, therapeutic pressure requirements when PAP is
required to effectively open or pneumatically splint a patient's upper airway.

25 With the two-piece, separate component design, as taught by Thornton, there is
a significant space between the upper and lower arches. Unfortunately, this
allows for PAP to escape at night when the patient is asleep which results in
lowering of therapeutic pressure. Attempts to correct this problem with prior art
have included adding a chin strap and/or a separate, extra component referred to
as a "vent or saliva shield." This vent shield must be added to the multi-
component device complicating utilization by patients.

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The present invention has no need for a chin strap or a separate vent shield to prevent leakage of requisite PAP. This problem is solved by sealing the upper and lower arches with acrylic to form a PAP obturator. There is no unsealed space between the upper and lower components anteriorly of the airway orthotic from which PAP can escape. Some space posteriorly may be intentionally left open for tongue room without compromising obturation because the tongue will fill the void laterally and complete the obturation. The snap fit of the teeth produced by the elastomeric material obviates the need for a chin strap or similar means to keep the mouth closed at night when PAP is blown into the nasal passages.

Advancing the mandible to correct OSAS can result in untoward occlusal changes or changes in the patient's bite. This occurs from significant orthodontic forces being applied to teeth due to intentional forward positioning of the mandible by using the maxillary teeth as an anchor or fulcrum to lever the mandible forward which results in muscles applying undesirable forces upon teeth at night.

The present invention solves this problem by placing the patient's lower jaw in a **neutral centric** position to minimize or eliminate orthodontic forces on the teeth where there is clinical indication to do so. However, the present invention may utilize mandibular advancement in the design when there may be some benefit to do so. The only reason to advance the mandible in combination therapy would be an attempt to reduce unreasonably high therapeutic PAP pressure if this factor is reducing patient compliance on nasal CPAP.

Means of application of PAP via the Thornton (combination) invention referred to as the "TAP/SAAMS" uses a one size fits all noses, tubing retention platform (SAAMS). In clinical practice this is very problematic as nose and nare widths vary widely. Without adjusting the width laterally of the PAP Tubing Retention Platform, many patients will fail this type of combination therapy. Thornton's art

does not anticipate the use of variable nose-width tubing retention platforms, nor is it obvious to those skilled in the art.

The present invention incorporates a method whereby the PAP Tubing Retention Platform is varied in lateral width to accompany the varied width of patient noses and nares. Different sizes in lateral widths of PAP Tubing Retention Platforms permit proper angulation and placement of nasal tubing carrying PAP. Proper angulation and placement of PAP Tubing results in a profound and stabile seal of PAP at the nares.

The present invention seeks to overcome prior art design that is fraught with another very significant problem as concerns proper angulation of nasal tubing to deliver PAP. This problem is that often the SAAMS platform is simply too far away, anterior-posteriorly, from the patient's nasal passages to approach the nares from a preferred angle. This is because the TAP/SAAMS design attaches or fits directly over the "Front Assembly (Part # CA-9001)" utilizing a Sheath Slide Extension. The anterior end of the Front Assembly, where the Adjustable Advancement Nut lies, is normally at least 22mm from the labial surface of the maxillary anterior teeth. When the SAAMS tubing retention platform is added/screwed into place on top of the Front Assembly via the Sheath Extension, the PAP nasal tubing is at least 35mm from the labial surface of the maxillary anterior teeth. This design flaw forces the clinician to angulate the nasal tubing in an undesirable manner often resulting in failure to properly seal the nares and maintain the seal while the patient is asleep.

The present invention allows the PAP Tubing Retention Platform to slide posteriorly to preferred positions which are much closer to the patient's nares. This allows for preferred positioning and angulation of PAP nasal tubing thereby improving the requisite seal at the nares and thereby increasing success in treatment. The present invention allows for the PAP Tubing Retention Platform

to achieve proximity to the labial aspect of the maxillary anterior teeth in a range of 5mm to 30mm depending on the patient's anatomy.

In addition to moving the PAP Tubing Retention Platform posteriorly on the
5 Variable Length Slide to achieve proximity of PAP Tubing directly below the
nares, the present invention recognizes additional means to assist in solving this
problem. First, the PAP Tubing Retention Platform may be "U" shaped at the
posterior end of the Platform such that the PAP Tubing Holes swing backward on
the left and right sides toward the patient's face. Second, the adjustable Slide
10 Mount Bracket may be mounted anterior to the posterior edge of the PAP Tubing
Retention Platform allowing the Platform to get closer to the patient's face.

The present invention also envisions a method of changing the Thornton design
whereby this limitation in design, which prevents close approximation of the PAP
15 Tubing Retention Platform to more closely approximate access to the nares, is
overcome. By removing the Adjustable Advancer Nut and inserting a female
threaded stainless steel sleeve into the posterior section of the Front Assembly, a
new Sheath Extension can be fit over this remaining section of the Front
Assembly to bring a SAAMS' PAP Tubing retention platform closer to the
20 preferred position underneath the patient's nares. Alternatively, a new Front
Assembly could be designed without the Adjustable Advancer Nut that includes a
built-in PAP Tubing Retention Slide that could be directly screwed into the
Locator Plate (Part # CA-9005) allowing closer approximation of the PAP Tubing
Retention Platform posteriorly. This would allow the current TAP/SAAMS user to
25 add PAP and reduce distance from the nares by at least 8mm. This would
significantly improve the combination therapy described by Thornton.

The "TAP/SAAMS" design incorporates a metal component termed the "Locator
Plate (Part # CA-9005)" which is imbedded within the anterior shell of acrylic in
30 the upper arch component of the TAP device. The Front Assembly with
Adjustable Advancer Nut is screwed into this Locator Plate via the Locator Nut

Set (Part # CA-9007). On top of this is the platform Sheath Extension that in turn screws onto the end of the Adjustable Advancer Nut. Because the Locator Plate is imbedded into the acrylic it can sustain excessive forces due to levering action of the Sheath Extension Slide such that there is a tendency for fracture of the acrylic at the anterior surface of the upper component of the MAD.

The present invention has no metal parts imbedded within the acrylic of the airway orthotic. The PAP Tubing Retention Platform Slide is bonded, injection molded, or cast in acrylic and thereby attached to the single piece, dual arch obturating airway orthotic. All exterior components are made from acrylic and bonded together. This greatly reduces potential for fracture. Additionally, in the present invention the length of the PAP Retention Platform Slide may be cut once the patient is fitted thereby reducing lever arm forces. This is not possible with the Thornton invention without irreparably damaging it.

This invention also seeks to improve on another nasal insert/pillow pressure application structure device known as the "Adam Circuit." The Adam Circuit is a headgear device sold by Puritan-Bennett Corp. (Pleasanton, CA) under the trademark "Breeze" that attempts to stabilize the use of nasal pillows. However, this device described in U.S. Patent 4,782,832 utilizes an uncomfortable and cumbersome headgear to support the sealing of the patient's nares with PAP. This device has a significant tendency to become displaced when a patient moves around in his bed at night or simply shift to his side. When this device is in contact with his bed-pillow the seal at the nares is often broken thereby reducing the requisite therapeutic pressure prescribed to open the patient's upper airway. The present invention utilizes nasal pillows but improves on the method of stabilization of the seal of the nares and avoids the uncomfortable and cumbersome headgear.

In reference to **U.S. Pat nts 5,752,510 and 6,012,455 (Goldstein)** there are novel and useful differences between the prior art and the present invention.

Goldstein describes an oral appliance that produces a double bite and uses PAP in combination. But there is no intent to obturate the oral cavity and the reason for Goldstein's dual arch is for a different purpose and his method of design and manufacture is quite clearly different than the present invention. The astute reader will note that the differences in the present invention are novel and useful as compared to Goldstein.

Description of Figures

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Figure 1 shows the present invention utilizing a single-piece, dual arch, obturator with an acrylically-connected slide upon which moves an adjustable PAP Tubing Retention Platform device.

15 Figure 2 shows the PAP Tubing Retention Platforms in variable widths (small, medium, and large) corresponding with variable nare and nasal widths.

Figure 3 shows a superior view of the dual arch, obturating airway orthotic with anterior supported for an array of varying width PAP Tubing Retention Platforms.

20 Only one size PAP Tubing Retention Platform is shown in this figure.

Description of the Preferred Embodiment

Figure 1 shows a lateral view of the present invention. This view is from the patient's right side where the Upper Right Quadrant #1 is superior and the Lower Right Quadrant #2 is inferior. Posteriorly there is Tongue Space #3 just where the Solid Acrylic Obturating Seal #8 ends. Typically the Solid Acrylic seal will extend to at least the mesial aspect of the maxillary first molar if not slightly further to create the most robust seal. The Upper Dental Arch #5 is superior to the Lower Dental Arch #6. The airway orthotic is composed of Exterior Hard Acrylic #7 and is lined interiorly with Elastomeric Material #4. Anteriorly the 5mm

Slide Mount #16 is operatively connected to the Variable Length Slide #13. Upon the Variable Length Slide the Variable Width PAP Tubing Retention Platform #12 is adjustably affixed to the Variable Length Slide via the Slide Mount Bracket #14. Superiorly and anteriorly there is a Nasal Pillow #9 which fits over a Collar #10.

- 5 The Collar inserts into the PAP Tubing #15 which is threaded through the Variable Width PAP Tubing Retention Platform via the PAP Tubing Hole (Figure 3, #18). At a certain preferred distance from the nose inferiorly is a Whisper Swivel II Valve (Respironics Part Number 332113). Above the Whisper Swivel II Valve and within the PAP Tubing is located an optional Exhaust Port #11. Both
10 the Whisper Swivel II Valve and the Exhaust Port are utilized to blow off or vent Carbon Dioxide (CO₂) on patient exhalation. The rate of expiratory flow from use of these structures is between 5-15 liters per minute. The need for the optional Exhaust Port can be determined in the sleep laboratory by end tidal CO₂ monitoring. If the patient is retaining too much CO₂ PAP Tubing Holes are
15 created to assist the Whisper Swivel II Valve in blowing off excess CO₂.

- Figure 2 shows the Present Invention whereby a method of treating Obstructive Sleep Apnea Syndrome utilizes PAP Tubing Retention Platforms that are variable in widths to correspond with variations in nasal and nare width. Figure 2
20 shows three hypothetical patient nasal widths small, medium, and large that correspond with particular width PAP Tubing Retention Platforms (small #12a, medium #12b, and large #12c). The reader will note that there could be additional Platforms as necessary to accommodate any and all variations in patient nasal widths. If there is flaring of the nares superiorly the PAP Tubing
25 Retention Platforms may be increased slightly in width to allow for angulation medially of the Platform via application of heat to bend the acrylic so as to customize the angular approach of the PAP Tubing #15. The preferred thickness of the Variable Width PAP Tubing Retention Platforms should be at least 3mm. These PAP Tubing Retention Platforms can be manufactured via standard
30 methods such as heat "suck-down" whereby a 3mm sheet of acrylic such as Biocryl is heated until pliable and vacuum-formed onto various width molds

forming the preferred structures. They may also be injection molded or cast directly in acrylic via the lost wax technique. The PAP Tubing Retention Platform is mounted onto the Slide #13 via the Slide Mount (Figure 1) and can be preferably located or adjusted as close as 5mm from the labial surface of the anterior teeth. The preferred range of location of the PAP Tubing #15 will be 5mm to 30mm from the labial surface of the maxillary anterior teeth. Once the preferred location antero-posteriorly is located for the individual patient the Slide #13 may be cut to reduce unnecessary lever arm forces.

Figure 3 shows the present invention in a superior view. The Variable Width PAP Tubing Retention Platform #12 is shown in two widths and styles in this figure and is mounted onto the Variable Length Slide #13 via the Slide Mount Bracket #14. The first style is rectangular and the second style is "U" shaped. There are two PAP Tubing Holes #18 which correspond to the right and left nares in both styles. The width of the PAP Tubing Retention Platform and position of the holes is preferably selected to match the corresponding width of the patient's nasal and nare width. The Solid Acrylic Obturating Seal #8 continues from the anterior region of the airway orthotic posteriorly as necessary to seal off the oral cavity from loss of requisite therapeutic positive airway pressure as determined in the sleep laboratory. There is Tongue Space #3 anteriorly, laterally, and vertically. The vertical or caudal dimension of the airway orthotic is varied according to tongue size and is determined by the experienced clinician through a variety of means or methods. The preferred embodiment utilizes a neuromuscular TENS (Transcutaneous Electrical Nerve Stimulation) technique whereby the masticatory muscles are profoundly relaxed to proper working lengths via this pulsing technique placing the mandible in three dimensional harmonious space position with respect to the maxilla. In this manner the mandibular position antero-posteriorly (AP) and vertically is determined by the muscles themselves rather setting an arbitrary position. This neuromuscularly-determined position is referred to earlier in the specification as "neutral centric" position. If the need presents, due to excessive therapeutic PAP

requirements for an individual patient, the AP position of the mandible may be somewhat protruded forward so as to create some mechanical dilation of the upper airway. However, this forward positioning of the mandible will increase the risk of a deleterious change in the patient's occlusion or bite. Therefore, the preferred position will typically be the neutral centric position as determined by the experienced clinician.

The foregoing description of the preferred embodiments of the invention has been presented for the purposes of illustration and description. It is not intended to be exhaustive or to limit the invention to the precise form disclosed. Accordingly, the foregoing description should be regarded as illustrative rather than restrictive, and it should be appreciated that variations may be made in the embodiments described by workers skilled in the art without departing from the scope of the present invention as defined by the following claims.

What is claimed is:

1) A method for treating Obstructive Sleep Apnea Syndrome (OSAS) with positive airway pressure (PAP) comprising:

- a) creating a single-piece, dual arch airway orthotic;
- b) using said airway orthotic to obturate the oral cavity via an acrylic seal between the upper and lower dental arches;
- c) retaining the upper and lower dental arches in elastomeric material via a snap-fit;
- d) applying PAP via the nasal passages from tubing which is supported from said airway orthotic.

2) The method of Claim 1) and additionally comprising a single-piece, dual arch airway orthotic whereby the lower (mandibular) arch is preferably located or placed into neutral centric position with respect to the maxilla.

3) The method of Claim 1) and additionally comprising utilization of variable width PAP Tubing Retention Platforms which correlate with varied patient nare and/or nasal widths.

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4) The method of Claim 1) and additionally comprising utilization of nasal pillows or inserts which are operably connected to said tubing.

5) The method of Claim 3) and additionally comprising customization of PAP
10 Tubing angulation by heating the PAP Tubing Retention Platform.

6) The method of Claim 5) where the means of heating is via a micro torch.

7) The method of Claim 1) and additionally comprising professional laboratory
15 processing of the single-piece, dual arch airway orthotic.

8) The method of Claim 7) and additionally comprising utilization of a three-dimensional bite registration.

20 9) The method of Claim 8) and additionally utilizing a three-dimensional bite registration that is created by a neuromuscular technique such as Transcutaneous Electrical Nerve Stimulation (TENS).

10) The method of Claim 8) whereby the three-dimensional bite registration
25 captures the mandibular position with respect to the maxilla in neutral centric.

11) A device which obturates the oral cavity during treatment of OSAS consisting of:

30 a) Single piece, dual arch oral appliance with a hard acrylic exterior and an interior lined with an elastomeric material;

b) Acrylically-bonded Variable Length Slide which supports a PAP Tubing Retention Platform that can vary the width of PAP Tubing Holes to correspond to varying nasal and nare widths.

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12) The device of Claim 11) and additionally comprising alignment of the upper and lower dental arches into a preferred position within said elastomeric material whereby the mandible is not protruded and is located in neutral centric.

10 13) The device in Claim 11) wherein said single piece, dual arch oral appliance has the upper and lower arch components sealed with hard acrylic.

14) The device of Claim 11) and additionally comprising alignment of the upper and lower dental arches into a preferred position within said elastomeric material
15 whereby the mandible is located in an anterior protruded position.

15) The device of Claim 12) and additionally comprising a means to create said preferred position of the mandible utilizing Transcutaneous Electrical Nerve Stimulation (TENS).

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16) The device of Claim 11) whereby said PAP Tubing Retention Platform can be positioned on said Variable Length Slide such that PAP Tubing can be positioned antero-posteriorly within a range of 5mm to 30mm from the labial surface of the maxillary anterior teeth.

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17) The device of Claim 11) whereby said variable width PAP Tubing Retention Platform is composed of acrylic material that is at least 3mm thick.

18) The method of applying PAP to nasal passages for the purpose of treating
30 OSAS comprising:

- a) Fabricating an oral appliance which has an anterior, extraoral Slide affixed to said oral appliance;
- b) Mounting a PAP Tubing Retention Platform to said extraoral Slide;
- c) Varying the lateral width of the PAP Tubing Holes in said PAP Tubing Retention Platform to correspond with varying width noses and nares.

19) The method of Claim 18) and further comprising positioning of said PAP Tubing Retention Platform and said PAP Tubing Holes antero-posteriorly to a position within a range of 5mm to 30mm from the labial surface of the maxillary anterior teeth.

20) The method of Claim 18) wherein said PAP Tubing Retention Platform is composed of acrylic material that is at least 3mm thick.

21) The method of Claim 20) wherein said acrylic material can be adjusted to optimize desired angulation via application of heat.

22) The method of Claim 18) wherein said PAP Tubing Retention Platform is vacuum formed subsequent to heating of the material.

23) The method of Claim 18) wherein said PAP Tubing Retention Platform is create via injection molding.

24) The method of Claim 18) wherein said oral appliance is a single-piece, dual arch obturator.

25) The method of Claim 24) where said obturator is composed of an exterior hard acrylic lined with an elastomeric material.

26) The method of Claim 24) where the position of the mandible within said single-piece, dual arch obturator is in neutral centric.

27) The method of Claim 24) where the position of the mandible within said single-piece, dual arch obturator is in forward protruded position.

5 28) The method of Claim 18) wherein said Variable Length Slide is acrylically bonded to the anterior surface of said oral appliance without the use of metal parts.

29) The method of Claim 18) where said oral appliance is composed of a hard
10 exterior acrylic and lined with an elastomeric material.

30) The method of Claim 18) where said oral appliance is manufactured in a professional laboratory.

15 31) The method of Claim 30) where fabrication of said oral appliance utilizes a three-dimensional bite registration to orient the position of the upper and lower dental arches.

32) The method of Claim 31) where said bite registration is produced utilizing
20 Transcutaneous Electrical Nerve Stimulation (TENS).

33) The method of treating OSAS utilizing a dual arch oral appliance without protrusion of the mandible such that the mandibular arch is located in neutral centric position with respect to the maxillary arch.

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34) The method of Claim 33) and further comprising support and stabilization of PAP Tubing from said dual arch oral appliance.

35) The method of Claim 34) and additionally comprising location of said PAP
30 Tubing through variable width PAP Tubing Retention Platforms whereby said PAP Tubing Holes correspond to varying nasal and nare widths.

36) The method of Claim 33) and additionally comprising capture of said neutral centric position via Transcutaneous Electrical Nerve Stimulation (TENS).

- 5 37) The method of Claim 33) and additionally comprising capture of said neutral centric position via conventional techniques such as manual physical manipulation of the mandible by the clinician.

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